

GenCore version 5.1.3  
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OM protein - protein search, using sw model

Run on: December 5, 2002, 13:18:38 ; Search time 38 Seconds

(without alignments)  
1171.203 Million cell updates/sec

Title: US-09-765-034-2

Perfect score: 1747  
Sequence: 1 MGIWAWATCKWMLAENAA.....KSLTSPRWAEHLTLSPREK 334

Scoring table: BLOSUM62  
Gapop 10.0 , Gapext 0.5

Searched: 908470 seqs, 133250620 residues

Total number of hits satisfying chosen parameters: 908470

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%  
Listing first 45 summaries

Database :

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2: /SID2/gcgdata/geneseq/geneseq-emb1/AA1981.DAT:\*  
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	1737	99.4	334	21	AAV71308 Human orphan G pro
2	1737	99.4	334	21	AA802842 Human G protein co
3	1737	99.4	334	23	AB890381 Human polypeptide
4	1737	99.4	379	23	AAE15633 Human G-protein co
5	1725	98.7	334	18	AAW19854 Human purinergic r
6	1721	98.5	334	18	AAW22732 Human ATP receptor
7	1696	97.1	387	22	AAU31029 Novel human secret
8	1533	76.3	258	21	AA845176 Human secreted pro
9	1231.5	70.5	317	23	AAU74904 Amino acid sequenc
10	499.5	28.6	373	23	AAU10984 Purinergic recepto

11	497.5	28.5	373	22	AAE04389	Human P2-purinergi
12	497.5	28.5	373	23	AAU10983	Purinergic recepto
13	490.5	28.1	373	23	AAU10985	Purinergic recepto
14	477	27.3	337	22	AAU04375	Human G-protein co
15	477	27.3	337	22	AAO15399	Human G-protein co
16	477	27.3	337	23	ABE81902	Human G-protein co
17	477	27.3	337	23	ABE83819	Human P21-like rec
18	477	27.3	337	23	ABE21803	Human AXOR89 (G-pr
19	477	27.3	337	23	ABE79438	Human p2y1-11. Ho
20	477	27.3	337	23	AAU77600	Human p2y1-like G
21	477	27.3	337	23	AAO14027	Human purinergic-r
22	477	27.3	337	23	AAE16171	Human G-protein co
23	475	27.2	374	22	AAE04390	Turkey P2Y nucleot
24	470	26.9	337	22	AAU04584	Human G-protein co
25	429.5	24.6	259	21	AA845375	Gene 37 human secr
26	420.5	24.1	336	22	AAE80971	Human NGPCRS4 #2.
27	405	23.2	537	23	AAU74538	Human P2Y purinoc
28	395.5	22.6	276	23	ABE83818	Human P2Y-1-like rec
29	386	22.1	373	23	AAE20604	Mus musculus GPCR
30	385	22.0	377	22	AAE04392	Human P2-purinergi
31	385	22.0	377	22	AAE01143	Human purinergic r
32	385	22.0	377	22	AAE01144	Human purinergic r
33	384	22.0	330	23	AAE77964	Human G-protein co
34	384	22.0	341	22	AAE07539	Human G-protein co
35	384	22.0	346	22	AAE12022	Human G-protein co
36	384	22.0	346	22	AAE82852	Human P2Y-1-like GPC
37	384	22.0	346	22	AAE07538	Human G-protein co
38	384	22.0	346	22	AAU04368	Human G-protein co
39	384	22.0	346	22	AAU07294	Cytosinyl leukorri
40	384	22.0	346	22	AAE73097	Human LTR4 recepto
41	384	22.0	346	22	ABE66684	Human novel polype
42	384	22.0	346	23	AAU10004	Human CysLT2-1-like
43	384	22.0	346	23	AAE17231	Human CysLT2 GPCR
44	384	22.0	346	23	AAE77965	Human G-protein co
45	384	22.0	346	23	ABE05229	Human LTD4-like G

#### ALIGNMENTS

RESULT 1	AAV71308	AAV71308 standard; Protein; 334 AA.
ID	AAV71308	
XX	AAV71308;	
AC	02-NOV-2000 (first entry)	
XX		
DT		
XX		
DE	Human orphan G protein-coupled receptor hCHN10.	
XX		
KW	Human; orphan G protein-coupled receptor; GPCR; hCHN10; drug screening;	
KW	Transmembrane Receptor; expressed sequence tag; EST; signal cascade.	
XX		
OS	Homo sapiens.	
XX		
PN	WO200031258-A2.	
XX		
PD	02-JUN-2000.	
XX		
PF	13-OCT-1999;	99WO-US23687.
XX		
PR	20-NOV-1998;	98US-0109213.
PR	16-FEB-1999;	99US-0120416.
PR	26-FEB-1999;	99US-0121852.
PR	12-MAR-1999;	99US-0123946.
PR	12-MAR-1999;	99US-0123949.
PR	28-MAY-1999;	99US-0136436.
PR	28-MAY-1999;	99US-0136437.
PR	28-MAY-1999;	99US-0136439.
PR	28-MAY-1999;	99US-0136567.
PR	28-MAY-1999;	99US-0137127.
PR	28-MAY-1999;	99US-0137131.
PR	29-JUN-1999;	99US-0141448.

PR 29-SEP-1999; 99US-0156555.  
 PR 29-SEP-1999; 99US-0156633.  
 PR 29-SEP-1999; 99US-0156634.  
 PR 29-SEP-1999; 99US-0156653.  
 PR 01-OCT-1999; 99US-0157280.  
 PR 01-OCT-1999; 99US-0157281.  
 PR 01-OCT-1999; 99US-0157282.  
 PR 01-OCT-1999; 99US-0157293.  
 PR 01-OCT-1999; 99US-0157294.  
 PR 12-OCT-1999; 99US-0416760.  
 PR 12-OCT-1999; 99US-0417044.  
 XX (AREN-) ARENA PHARM INC.  
 XX  
 XX Chen R, Dang HT, Liaw CW, Lin I;  
 XX  
 XX WPI; 2000-400068/34.  
 DR N-PSDB; AAD01135.  
 XX  
 XX Novel human orphan G protein-coupled receptors and the encoding cDNAs  
 PT for use in the identification of G protein-coupled receptor agonists -  
 XX  
 XX Claim 70; Page 87-88; 102pp; English.  
 XX  
 CC The present amino acid sequence is the hCHN10, an endogenous human  
 CC orphan G protein-coupled receptor (GPCR), expressed in kidney and  
 CC thyroid. The hCHN10 cDNA was identified using the human EST (expressed  
 CC sequence tag) 1365839 as a probe. The orphan GPCR of the invention, like  
 CC all GPCRs has seven transmembrane alpha helices with an extracellular  
 CC N-terminus and an intracellular C-terminus. However, no endogenous  
 CC ligands has yet been identified for the proteins of the invention. The  
 CC orphan GPCRs may be used in the identification of their endogenous  
 CC ligands, and to screen potential GPCR agonists and antagonists for use as  
 CC pharmaceutical agents. The proteins may also be used in the study of  
 CC GPCR-mediated signalling cascades, and to elucidate their precise role in  
 CC normal and diseased human conditions. Nucleic acid encoding human orphan  
 CC GPCRs may be used for tissue localisation expression analysis to provide  
 CC information about their function in healthy and pathological states.  
 XX  
 XX Sequence 334 AA;  
 SQ  
 Query Match 99.4%; Score 1737; DB 21; Length 334;  
 Best Local Similarity 99.7%; Pred. No. 8.2e-172;  
 Matches 333; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
 QY 1 MGIAMNATCKNWLAAEAALEKYLSIFGYIEFVGVGLNTIVVGYIFSLKWNSSNI 60  
 Db 1 MGIAMNATCKNWLAAEAALEKYLSIFGYIEFVGVGLNTIVVGYIFSLKWNSSNI 60  
 QY 61 YLFNLSVSDLAFLCTPLMIRSYANGNIYGDVLCISNRYVLANLYTSILFLTFISDR 120  
 Db 61 YLFNLSVSDLAFLCTPLMIRSYANGNIYGDVLCISNRYVLANLYTSILFLTFISDR 120  
 QY 121 YLIITKYPREHLLQKFAILISIAVWLTLELLPLINPVIITNGTTCNDFASSGD 180  
 Db 121 YLIITKYPREHLLQKFAILISIAVWLTLELLPLINPVIITNGTTCNDFASSGD 180  
 QY 181 PNYNLISYMCULTLGLLPIPLVMCFYFKIALFLKQRNRQVATLPLEKPLNLVIMAVVI 240  
 Db 181 PNYNLISYMCULTLGLLPIPLVMCFYFKIALFLKQRNRQVATLPLEKPLNLVIMAVVI 240  
 QY 241 FSVPTPHVMNRVIRASRLGSKQYQCTQVINSFIVTRPLAFLNSVNPVYFLFGD 300  
 Db 241 FSVPTPHVMNRVIRASRLGSKQYQCTQVINSFIVTRPLAFLNSVNPVYFLFGD 300  
 QY 301 HFRDMLNQLRHNFKSLTSFSRWAEHLLSPREK 334  
 Db 301 HFRDMLNQLRHNFKSLTSFSRWAEHLLSPREK 334  
 RESULT 2  
 AAB02842  
 ID AAB02842 standard; Protein; 334 AA.

XX AAB02842;  
 XX 22-AUG-2000 (first entry)  
 XX Human G protein coupled receptor hCHN10 protein SEQ ID NO:38.  
 XX Human; G protein coupled receptor; GPCR; transmembrane receptor;  
 KW identification; agonist; screening; therapeutic; pharmaceutical;  
 XX mutant.  
 XX Homo sapiens.  
 XX WO200022131-A2.  
 XX 20-APR-2000.  
 XX 13-OCT-1999; 99WO-US24065.  
 XX 13-OCT-1998; 98US-0170496.  
 PR 12-NOV-1998; 98US-0108029.  
 PR 20-NOV-1998; 98US-0109213.  
 PR 27-NOV-1998; 98US-0110060.  
 PR 16-FEB-1999; 99US-0120416.  
 PR 26-FEB-1999; 99US-0121852.  
 PR 12-MAR-1999; 99US-0123944.  
 PR 12-MAR-1999; 99US-0123945.  
 PR 12-MAR-1999; 99US-0123946.  
 PR 12-MAR-1999; 99US-0123948.  
 PR 12-MAR-1999; 99US-0123949.  
 PR 12-MAR-1999; 99US-0123951.  
 PR 28-MAY-1999; 99US-0136436.  
 PR 28-MAY-1999; 99US-0136437.  
 PR 28-MAY-1999; 99US-0136439.  
 PR 28-MAY-1999; 99US-0137127.  
 PR 28-MAY-1999; 99US-0137131.  
 PR 28-MAY-1999; 99US-0137567.  
 PR 30-JUN-1999; 99US-0141448.  
 PR 27-AUG-1999; 99US-0151114.  
 PR 03-SEP-1999; 99US-0152524.  
 PR 29-SEP-1999; 99US-0156633.  
 PR 29-SEP-1999; 99US-0156555.  
 PR 29-SEP-1999; 99US-0156634.  
 XX (AREN-) ARENA PHARM INC.  
 PA Behan DP, Lehmann-Bruinsma K, Chalmers DT, Chen R, Dang HT;  
 PI Gore M, Liaw CW, Lin I, Lowitz K, White C;  
 XX WPI; 2000-317986/27.  
 DR N-PSDB; AAA46036.  
 XX Non-endogenous, human G protein-coupled receptors for screening  
 PT receptor, inverse or partial agonists useful as therapeutic agents -  
 XX Example 1; Page 117-118; 187pp; English.  
 PS The present invention describes transmembrane receptors, preferably  
 CC human G protein coupled receptors (GPCR), for which the endogenous  
 CC ligand is unknown (Orphan GPCR receptors). More specifically the present  
 CC invention relates to non-endogenous, constitutively activated versions  
 CC of a human GPCR. These non-endogenous human GPCRs can be useful for  
 CC the direct identification of candidate compounds as receptors agonists,  
 CC inverse agonists or partial agonists for use as pharmaceutical agents.  
 CC AAA46017 to AAA46126 and AAB02825 to AAB02859 represent sequences used in  
 CC the exemplification of the present invention.  
 XX Sequence 334 AA;  
 SQ  
 Query Match 99.4%; Score 1737; DB 21; Length 334;  
 Best Local Similarity 99.7%; Pred. No. 8.2e-172;  
 Matches 333; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 MGIMAMNATCKNWLAAEALAEKYYLSTFYGIEFVVGVTGNTIVVGYIFSLKNNSSNI 60  
 DB 1 MGIMAMNATCKNWLAAEALAEKYYLSTFYGIEFVVGVTGNTIVVGYIFSLKNNSSNI 60  
 QY 61 YFNLVSVDLAFCLTLPMLIRSYANGNWIYGDVLCISNRVYVLANLYTSLFLTFISIDR 120  
 DB 61 YFNLVSVDLAFCLTLPMLIRSYANGNWIYGDVLCISNRVYVLANLYTSLFLTFISIDR 120  
 QY 121 YLIKKYPRFHHLLQKKEFALISLAIWVLTLELPLPLINPVITDNGTTCNDPSSGD 180  
 DB 121 YLIKKYPRFHHLLQKKEFALISLAIWVLTLELPLPLINPVITDNGTTCNDPSSGD 180  
 QY 181 PNYNLIYSMCLTLGLFLPLFVWCFFYYKIALFLKQRNQVATALPLEKPLNLVIAVVI 240  
 DB 181 PNYNLIYSMCLTLGLFLPLFVWCFFYYKIALFLKQRNQVATALPLEKPLNLVIAVVI 240  
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 DB 241 FSVLPFTPYHMRNVRIRASLGSWKQYQCTOVVINSFYITRPLAFINSYINPVFYFLGD 300  
 QY 301 HPRDMLMNLQRHNFKSLTFSRMAHLLSFRK 334  
 DB 301 HPRDMLMNLQRHNFKSLTFSRMAHLLSFRK 334

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 ID ABB90381 standard; Protein; 334 AA.  
 XX ABB90381;  
 AC ABB90381;  
 XX 24-MAY-2002 (first entry)  
 DT 24-MAY-2002 (first entry)  
 XX Human polypeptide SEQ ID NO 2757.  
 DE Human polypeptide SEQ ID NO 2757.  
 XX Cytostatic; immunosuppressive; nootropic; neuroprotective; antiviral;  
 KW antiallergic; hepatotropic; antidiabetic; antiinflammatory; antitumor;  
 KW vulnery; anticonvulsant; antibacterial; antifungal; antiparasitic;  
 KW cardiac; gene therapy; cancer; immune disorder; cardiovascular disorder;  
 KW neurological disease; infection; human; secreted protein.  
 XX Homo sapiens.  
 OS Homo sapiens.  
 XX WO200190304-A2.  
 PN WO200190304-A2.  
 XX 29-NOV-2001.  
 PD 29-NOV-2001.  
 XX 18-MAY-2001; 2001WO-US16450.  
 PF 18-MAY-2001; 2001WO-US16450.  
 XX 19-MAY-2000; 2000US-205515P.  
 PR 19-MAY-2000; 2000US-205515P.  
 XX (HUMA-) HUMAN GENOME SCI INC.  
 PA (HUMA-) HUMAN GENOME SCI INC.  
 XX Birse CE, Rosen CA;  
 PI Birse CE, Rosen CA;  
 XX WPI: 2002-122018/16.  
 DR N-PSDB; ABL90790.  
 XX Novel 1405 isolated polypeptides, useful for diagnosis, treatment and  
 PT prevention of neural, immune system, muscular, reproductive,  
 PT gastrointestinal, pulmonary, cardiovascular, renal and proliferative  
 PT disorders -  
 XX Claim 11; SEQ ID NO 2757; 2081bp + Sequence Listing; English.  
 PS The invention relates to novel genes (ABR9449-ABR90853) and proteins  
 CC (ABR9440-ABR90444) useful for preventing, treating or ameliorating  
 CC medical conditions e.g. by protein or gene therapy. The genes are  
 CC isolated from a range of human tissues disclosed in the specification.  
 CC The nucleic acids, proteins, antibodies and (ant)agonists are useful  
 CC in the diagnosis, treatment and prevention of: (a) cancer, e.g. breast  
 CC and ovarian cancer and other cancers of the adrenal gland, bone, bone  
 CC marrow, breast, gastrointestinal tract, liver, lung, or urogenital;

CC (b) immune disorders e.g. Addison's disease, allergies, autoimmune  
 CC haemolytic anaemia, autoimmune thyroiditis, diabetes mellitus, Crohn's  
 CC disease, multiple sclerosis, rheumatoid arthritis and ulcerative  
 CC colitis; (c) cardiovascular disorders such as myocardial ischaemia;  
 CC (d) wound healing; (e) neurological diseases e.g. cerebral anoxia and  
 CC epilepsy; and (f) infectious diseases such as viral, bacterial, fungal  
 CC and parasitic infections.  
 CC Note: The sequence data for this patent did not form part of the  
 CC printed specification, but was obtained in electronic format directly  
 CC from WIPO at ftp.wipo.int/pub/published\_pct\_sequences.

XX Sequence 334 AA:

Query Match 99.4%; Score 1737; DB 23; Length 334;

Best Local Similarity 99.7%; Pred. No. 8,2e-172;  
 Matches 333; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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 QY 61 YFNLVSVDLAFCLTLPMLIRSYANGNWIYGDVLCISNRVYVLANLYTSLFLTFISIDR 120  
 DB 61 YFNLVSVDLAFCLTLPMLIRSYANGNWIYGDVLCISNRVYVLANLYTSLFLTFISIDR 120  
 QY 121 YLIKKYPRFHHLLQKKEFALISLAIWVLTLELPLPLINPVITDNGTTCNDPSSGD 180  
 DB 121 YLIKKYPRFHHLLQKKEFALISLAIWVLTLELPLPLINPVITDNGTTCNDPSSGD 180  
 QY 181 PNYNLIYSMCLTLGLFLPLFVWCFFYYKIALFLKQRNQVATALPLEKPLNLVIAVVI 240  
 DB 181 PNYNLIYSMCLTLGLFLPLFVWCFFYYKIALFLKQRNQVATALPLEKPLNLVIAVVI 240  
 QY 241 FSVLPFTPYHMRNVRIRASLGSWKQYQCTOVVINSFYITRPLAFINSYINPVFYFLGD 300  
 DB 241 FSVLPFTPYHMRNVRIRASLGSWKQYQCTOVVINSFYITRPLAFINSYINPVFYFLGD 300  
 QY 301 HPRDMLMNLQRHNFKSLTFSRMAHLLSFRK 334  
 DB 301 HPRDMLMNLQRHNFKSLTFSRMAHLLSFRK 334

RESULT 4  
 AAE15633  
 ID AAE15633 standard; Protein; 379 AA.  
 XX AAE15633;  
 AC AAE15633;  
 XX 12-MAR-2002 (first entry)  
 DT 12-MAR-2002 (first entry)  
 XX Human G-protein coupled receptor-3 (GCR3-3) protein.  
 DE Human G-protein coupled receptor-3 (GCR3-3) protein.  
 XX Human; G-protein coupled receptor-3; GCR3-3; therapy; cancer; stroke;  
 KW cell proliferative disorder; neurological; epilepsy; Parkinson's disease;  
 KW Alzheimer's disease; inflammation; thyroiditis; haemolytic anaemia; AIDS;  
 KW Acquired Immune Deficiency Syndrome; dementia; nootropic; cholelithiasis;  
 KW multiple sclerosis; atherosclerosis; angina pectoris; gastroenteritis;  
 KW diabetes; ulcer; viral infection; immunosuppressive.  
 XX Homo sapiens.  
 OS Homo sapiens.  
 XX Key Location/Qualifiers  
 FH 187..206  
 FT Domain //label= Transmembrane\_domain  
 FT 234..253  
 FT Domain //label= Transmembrane\_domain  
 FT 276..296  
 FT Domain //label= Transmembrane\_domain  
 FT 319..342  
 FT Domain //label= Transmembrane\_domain  
 XX WO200190351-A2.



```
XX WPI; 1997-310601/28.
DR N-PSDB; AAT71900.
XX
XX New isolated purinergic receptor sub-type - used to develop
PT products for diagnosis and therapy, e.g. for screening for agonists
PT and antagonists which can modulate activation
XX
XX Claim 1; Fig 1A-B; 36pp; English.
XX
CC P2U2 receptor (AAW19854) is a novel human purinergic receptor
CC subtype that is abundantly expressed in kidney and in many cell
CC lines of megakaryocytic or erythroleukemic origin and which is
CC activated by ATP, UDP, UTP and dUDP. Its amino acid sequence was
CC deduced from a cDNA clone derived from DAMI (ATCC CRL 9792) cells.
CC P2U2 and its polypeptides can be expressed in host cells and used
CC to develop diagnostic and therapeutic agents. Antagonists and
CC agonists based on the extracellular domains of P2U2 receptor, or
CC which affect receptor function by binding to one of the
CC intracellular domains, can be used to treat diseases caused by
CC aberrant activation of this receptor or to treat diseases whose
CC symptoms can be ameliorated by stimulating or inhibiting the
CC activity of the receptor.
XX
SQ Sequence 334 AA;
XX
Query Match 98.7%; Score 1725; DB 18; Length 334;
Best Local Similarity 99.1%; Pred. No. 1.4e-170;
Matches 331; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
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QY 1 MGIAMWATCKNMLAABALEKYYLSIFYGIEFVGVGNTIVVGYIFSLKNNSSNI 60
DB 1 MGIAMWATCKNMLAABALEKYYLSIFYGIEFVGVGNTIVVGYIFSLKNNSSNI 60
XX
QY 61 YLFNLSVSDLAFLCTLPMLIRSYANGNMIYGDVLCISNRYVLANLYSILFLFTISIDR 120
DB 61 YLFNLSVSDLAFLCTLPMLIRSYANGNMIYGDVLCISNRYVLANLYSILFLFTISIDR 120
XX
QY 121 YLIIRYPRREHLQKKEFAIILISLAIWLVNTELELPILPINPVITDNGTCTNDPSSGD 180
DB 121 YLIIRYPRREHLQKKEFAIILISLAIWLVNTELELPILPINPVITDNGTCTNDPSSGD 180
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DB 241 FSVPTPYHVMNRVRIASRLGSMKQYQCTQVIVNSFYIVTRPALNSVINVEFYLLGD 300
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QY 301 HFRDMLNQLRHNFKSLTFSRMAHELLLSPREK 334
DB 301 HFRDMLNQLRHNFKSLTFSRMAHELLLSPREK 334
XX
RESULT 6
AAW22732
ID AAW22732 standard; Protein: 334 AA.
XX
AC AAW22732;
XX
DT 07-OCT-1997 (first entry)
XX
DE Human ATP receptor.
XX
KW ATP receptor; G-protein coupled receptor; agonist; antagonist.
XX
OS Homo sapiens.
XX
XX Key Location/Qualifiers
FH Misc-difference 212
FT /note= "encoded by TCC"
FT Misc-difference 235
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FT /note= "encoded by TCG"
FT Misc-difference 244
FT /label= Unknown
FT /note= "encoded by CYT"
XX
XX W09724929-A1.
XX
XX 17-JUL-1997.
XX
XX 11-JAN-1996; 96WO-US00392.
XX
XX 11-JAN-1996; 96WO-US00392.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Li Y;
XX
XX WPI; 1997-372505/34.
DR N-PSDB; AAT75146.
XX
XX Isolated human ATP receptor - agonists and antagonists of which are
PT useful in treatment of, e.g. asthma, hypertension, arterial
PT thrombosis and psychotic and neurological disorders
XX
XX Claim 15; Fig 1A-C; 53pp; English.
XX
XX Human ATP receptor (AAW22732) is structurally related to the G
XX protein-coupled receptor family. It shows 29.8% identity to a
XX murine P2u receptor. Its amino acid sequence was deduced from a
XX human placental cDNA clone (AAT75146). Recombinant ATP receptor can
XX be expressed in bacterial (e.g. E. coli), mammalian (e.g. COS) or
XX insect (e.g. Sf9) host cells and used to screen for agonists and
XX antagonists useful in the treatment of conditions related to
XX underexpression of the receptor (e.g. asthma, Parkinson's disease,
XX acute heart failure, hypotension, urinary retention and
XX osteoporosis) or underexpression of the receptor (e.g. arterial
XX thrombosis, hypertension, thrombolysis, angioplasty, cystic
XX fibrosis, ulcers, asthma, allergy, benign prostatic hypertrophy,
XX psychotic and neurological disorders, dyskinesias, endogenous
XX anorexia and bulimia).
XX
SQ Sequence 334 AA;
XX
Query Match 98.5%; Score 1721; DB 18; Length 334;
Best Local Similarity 99.5%; Pred. No. 3.7e-170;
Matches 329; Conservative 3; Mismatches 2; Indels 0; Gaps 0;
XX
QY 1 MGIAMWATCKNMLAABALEKYYLSIFYGIEFVGVGNTIVVGYIFSLKNNSSNI 60
DB 1 MGIAMWATCKNMLAABALEKYYLSIFYGIEFVGVGNTIVVGYIFSLKNNSSNI 60
XX
QY 61 YLFNLSVSDLAFLCTLPMLIRSYANGNMIYGDVLCISNRYVLANLYSILFLFTISIDR 120
DB 61 YLFNLSVSDLAFLCTLPMLIRSYANGNMIYGDVLCISNRYVLANLYSILFLFTISIDR 120
XX
QY 121 YLIIRYPRREHLQKKEFAIILISLAIWLVNTELELPILPINPVITDNGTCTNDPSSGD 180
DB 121 YLIIRYPRREHLQKKEFAIILISLAIWLVNTELELPILPINPVITDNGTCTNDPSSGD 180
XX
QY 181 PNYNLIYSMCLTLLGFLIPLFVMCFYKYIALFLKORNRQVATALPLEKPLNLVIMAVI 240
DB 181 PNYNLIYSMCLTLLGFLIPLFVMCFYKYIALFLKORNRQVATALPLEKPLNLVIMAVI 240
XX
QY 241 FSVPTPYHVMNRVRIASRLGSMKQYQCTQVIVNSFYIVTRPALNSVINVEFYLLGD 300
DB 241 FSVPTPYHVMNRVRIASRLGSMKQYQCTQVIVNSFYIVTRPALNSVINVEFYLLGD 300
XX
QY 301 HFRDMLNQLRHNFKSLTFSRMAHELLLSPREK 334
DB 301 HFRDMLNQLRHNFKSLTFSRMAHELLLSPREK 334
XX
RESULT 7
```

```
AAU31029
ID AAU31029 standard; Protein; 387 AA.
XX
AC AAU31029;
XX
DT 18-DEC-2001 (first entry)
XX
XX Novel human secreted protein #1520.
DE
XX Human; vaccination; gene therapy; nutritional supplement;
KW stem cell proliferation; haematopoiesis; nerve tissue regeneration;
KW immune suppression; immune stimulation; anti-inflammatory; leukaemia.
XX
XX Homo sapiens.
OS
XX WO200179449-A2.
XX
XX 25-OCT-2001.
XX
XX 16-APR-2001; 2001WO-US08656.
XX
XX 18-APR-2001; 2000US-0552929.
XX
XX 26-JAN-2001; 2001US-0770160.
XX
XX (HYSE-) HYSEQ INC.
XX
XX Tang YT, Liu C, Drmanac RT;
PI
XX WPI; 2001-611725/70.
XX
XX Nucleic acids encoding a range of human polypeptides, useful in genetic
XX vaccination, testing and therapy -
XX
XX Claim 20; Page 392; 765pp; English.
XX
XX The invention relates to novel human secreted polypeptides. The
XX polypeptides and antibodies to the polypeptides are useful for
XX determining the presence of or predisposition to a disease associated
XX with altered levels of polypeptide. The polypeptides are also useful for
XX identifying agents (agonists and antagonists) that bind to them. Cells
XX expressing the proteins are useful for identifying a therapeutic agent
XX for use in treatment of a pathology related to aberrant expression or
XX physiological interactions of the polypeptide. Vectors comprising
XX the nucleic acids encoding the polypeptides and cells genetically
XX engineered to express them are also useful for producing the proteins.
XX The proteins are useful in genetic vaccination, testing and
XX therapy, and can be used as nutritional supplements. They may be used to
XX increase stem cell proliferation; to regulate haematopoiesis; and in
XX bone, cartilage, tendon and/or nerve tissue growth or regeneration;
XX immune suppression and/or stimulation; as anti-inflammatory agents; and
XX in treatment of leukaemias. AAU29510-AAU3304 represent the amino acid
XX sequences of novel human secreted proteins of the invention.
XX
XX Sequence 387 AA;
XX
XX Query Match 97.1%; Score 1696; DB 22; Length 387;
XX Best Local Similarity 98.8%; Pred. No. 1.8e-167;
XX Matches 325; Conservative 1; Mismatches 3; Indels 0; Gaps 0;
XX
XX 6 AWWATCKWLAEEALEKYLISIFGIEFVGVGLGNTIVVGYIFSLKNWSSNLYLNL 65
XX |||||
XX 59 AWWATCKWLAEEALEKYLISIFGIEFVGVGLGNTIVVGYIFSLKNWSSNLYLNL 118
XX |||||
XX 66 SVSDLAFLCTPLMLRSVANGNWIQDVLCISNRVVLHANLYTSILFTFISIDRYLIK 125
XX |||||
XX 119 SVSDLAFLCTPLMLRSVANGNWIQDVLCISNRVVLHANLYTSILFTFISIDRYLIK 178
XX |||||
XX 126 YPFREHLLQKKEFAILISLAIWLVLTLELLPLILPINVITDGTGTCNDNFASSGDPNYNL 185
XX |||||
XX 179 YPFREHLLQKKEFAILISLAIWLVLTLELLPLILPINVITDGTGTCNDNFASSGDPNYNL 238
XX |||||
XX 186 IYSMCLTLLGFLPFLVCMCFYFKIALFLKQRNRQVATLPLEKPLNLVIMAVVIFSVLF 245
XX |||||
```

CC polypeptides can also be used as a food additive or preservative to  
CC increase or decrease storage capabilities, fat content, lipid, protein,  
CC carbohydrate, vitamins, minerals, cofactors and other nutritional  
CC components. AAC81077 to AAC81085 and AAB45307 represent sequences used in  
CC the exemplification of the present invention.

XX Sequence 258 AA;

Query Match 76.3%; Score 1333; DB 21; Length 258;  
Best Local Similarity 99.6%; Pred. No. 5,2e-130;  
Matches 257; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 24 YLLSIFVGIIEFVGVGNTIVVGYIFSLKMNSSNIYLFNLVSDFLCTPLMLIRSY 83  
DB 1 YLLSIFVGIIEFVGVGNTIVVGYIFSLKMNSSNIYLFNLVSDFLCTPLMLIRSY 60  
QY 84 ANGNWIVDVLCISNRVYLVHANLYTSILFTFISIDRYLIKPPREHLQKKEPAIILIS 143  
DB 61 ANGNWIVDVLCISNRVYLVHANLYTSILFTFISIDRYLIKPPREHLQKKEPAIILIS 120  
QY 144 LAIWLVTLELLPLILPLINPVITDNGTTCNDFASSGDPVYVNLISMCITLGLFLLPLFVM 203  
DB 121 LAIWLVTLELLPLILPLINPVITDNGTTCNDFASSGDPVYVNLISMCITLGLFLLPLFVM 180  
QY 204 CFYFYKIALFLKORNRQVATLPLEKPLNLVIMAVVIFSVPPTPYHVMNRVIAASRLGSM 263  
DB 181 CFYFYKIALFLKORNRQVATLPLEKPLNLVIMAVVIFSVPPTPYHVMNRVIAASRLGSM 240  
QY 264 KOYQCTQVIVNSFYIVTR 281  
DB 241 KOYQCTQVIVNSFYIVTR 258

## RESULT 9

AAU74904  
ID AAU74904 standard; Protein, 317 AA.

XX AAU74904;  
XX

DT 09-APR-2002 (first entry)

XX Amino acid sequence of mouse G-protein coupled receptor TGR18 protein.

XX Mouse; G-protein coupled; receptor; GPCR; TGR18; kidney disease;  
XX signal transduction modulator; cerebral cavernous malformation;  
XX hyperlipidemia; obesity; dyslexia; cardiac myxoma; renal failure;  
XX nephritis; hypertension; liver disease; cirrhosis; blood disorder;  
XX spleen-associated disorder; immune disorder.

XX Mus sp.

XX W0200200719-A2.

XX 03-JAN-2002.

XX 25-JUN-2001; 2001WO-US20363.

XX 23-JUN-2000; 2000US-213461P.

XX (TULA-) TULARIK INC.

XX Lin DC, Zhao J, Chen J, Cutler G;

XX WPI; 2002-147880/19.

XX N-PSDB; ABK12957.

XX New G-protein coupled receptor polypeptides, useful for identifying  
PT modulators of signal transduction for treating kidney disease,  
PT hyperlipidemia, obesity, dyslexia and cardiac myxoma -

XX Claim 33; Page 59; 78pp; English.

XX The present invention relates to a new G-protein coupled receptor (GPCR)

CC polypeptide comprising greater than 70% amino acid sequence identity to  
CC the amino acid sequence of human GPCRs TGR62, TGR21, TGR130.1, TGR130.2,  
CC human TGR23 or TGR92, 80% amino acid sequence identity to mouse TGR18  
CC or 90% amino acid sequence identity to human novel edg receptor protein,  
CC as defined in the specification. The GPCR covalently linked to a solid  
CC phase is useful for identifying a compound that modulates signal  
CC transduction. The identified compounds are useful for treating  
CC kidney disease, cerebral cavernous malformations, hyperlipidemia,  
CC obesity, dyslexia and cardiac myxoma. The molecules of the invention are  
CC useful for diagnosing disorders or conditions such as kidney-related  
CC conditions or diseases such as renal failure, nephritis, nephrotic  
CC syndrome, asymptomatic urinary abnormalities, renal tubule defects,  
CC hypertension and nephrolithiasis, liver-related disease or condition  
CC e.g. cirrhosis, infiltrations, lesions, functional disorders and jaundice  
CC and spleen-associated disorders or conditions e.g. splenic enlargement,  
CC immune disorders, blood disorders and others. Modulation of the  
CC polypeptide of the invention is useful to treat or prevent any of the  
CC above conditions or diseases. The present amino acid sequence represents  
CC the mouse GPCR TGR18 protein of the invention. This sequence is one of  
CC seven novel G protein coupled receptors of the invention (AAU74904-  
CC AAU74911).

XX Sequence 317 AA;

Query Match 70.5%; Score 1231.5; DB 23; Length 317;  
Best Local Similarity 71.8%; Pred. No. 2,4e-119;  
Matches 227; Conservative 42; Mismatches 46; Indels 1; Gaps 1;

QY 5 MAMNATCKNWLAEALAEKYYISIFVGIIEFVGVGNTIVVGYIFSLKMNSSNIYLFN 64  
DB 1 MAQNLSCEWNLATEAILNKYYLSAFYAIEFIFGLGNVTVVGYIFCMKMNSSNIYLFN 60

QY 65 LVSDFLFLCTPLMLIRSYANGNMYGDLCTSNRVYLVHANLYTSILFTFISIDRYLI 124  
DB 61 LVSDFLFLCTPLMLIRSYANGNMYGDLCTSNRVYLVHANLYTSILFTFISIDRYLI 120

QY 125 KYPPREHLQKKEPAIILISLAIWVLTLELLPLILPLINPVITDNGTTCNDFASSGDPVNY 184  
DB 121 KYPPREHLQKKEPAIILISLAIWVLTLELLPLILPLINPVITDNGTTCNDFASSGDPVNY 180

QY 165 LIYSNCLTLGLFLLPLFVWCFYFYKIALFLKORNRQVATLPLEKPLNLVIMAVVIFSVP 244  
DB 161 LIYSNCLTLGLFLLPLFVWCFYFYKIALFLKORNRQVATLPLEKPLNLVIMAVVIFSVP 240

QY 245 FPPYHVMNRVIAASRLGSMKOYQCTQVIVNSFYIVTRPLAFNSVINYVFFYLLGDHFD 304  
DB 241 FPPYHVMNRVIAASRLGSMKOYQCTQVIVNSFYIVTRPLAFNSVINYVFFYLLGDHFD 299

QY 305 MLMNQLRHNPKSLTSF 320  
DB 300 MLMNQLRHNPKSLTSF 315

## RESULT 10

AAU10984  
ID AAU10984 standard; Protein, 373 AA.

XX AAU10984;  
XX

DT 12-MAR-2002 (first entry)

XX Purinergic receptor P2Y, G-protein coupled 1, isoform #1.

XX Purinergic receptor P2Y, G-protein coupled 1; P2RY1; anticoagulant;  
XX coagulant; platelet aggregation; haplotyping; drug screening;

XX transgenic animal; human.

XX Homo sapiens.

XX Key Location/Qualifiers

XX Misc-difference 34 /note= "Wild type Ala substituted by Val"

XX FT





CC restenotic complications following angioplasty, carotid endarterectomy,  
CC post CABG (coronary artery bypass graft) surgery, vascular graft surgery,  
CC stent placements or insertion of endovascular devices and prostheses.  
CC P2Y12 receptor is useful for identifying binding partners and for  
CC diagnostic applications. P2Y12 receptor provides targets for screening  
CC synthetic small molecules and combinatorial or naturally occurring  
CC compound libraries to regulate platelet aggregation, vascular injury, or  
CC disease as well as schizophrenia, eating disorders, depression, migraine  
CC and other brain disorders. The present sequence is human P2-purinergic  
CC receptor subtype, P2Y1 related to the invention.

XX Sequence 373 AA;

Query Match 28.5%; Score 497.5; DB 22; Length 373;

Best Local Similarity 36.3%; Pred. No. 5e-43; Mismatches 124; Indels 21; Gaps 9;

Matches 117; Conservative 60; Mismatches 124; Indels 21; Gaps 9;

QY 6 AW-NATCNMMLAA---BALEK---YLSIFYGIEFVGVGLNTTVVGYIFSLKNW 55

DB 24 SWGNSVASTAAVSSSFICALTKTGFOFYIPLPAVYILVFIIIGFLNSVAIMVFFHMKRW 83

QY 56 NSSNIYLFNLSVSDIAFLCTLPMLIRSYAN-GNWIYGDVLCISNRVYLANLYTSILFLT 114

DB 84 SGISVYMFNMLADFLVLTLPALIFYYFNKTDWIFGDMCKLQRFIFHVNLVGSILFLT 143

QY 115 FISIDRYLIKYPREHLLQKKEFAILISLAIVWVLTLELPILPLINPVITDNGT-TCN 173

DB 144 CISAHRYSGVYVPLKSLRLKKKNAICISVLWLVVAISILYSGTGVRKNNTITCY 203

QY 174 DPASGDPNVLIIYSMCLTLGLFIPFVM--CFPYVKIALFLKQRNRQVATAPL-EKP 230

DB 204 DTSDEYLRSYFIYSMCTTVAMFCVPLVILIGCYGLIVRALYKDLNLS---PLRRKS 258

QY 231 LNLVIMAVIFSVPPTPYHVMNRVRIASRLGSKQYOCT-QVINSFYIVTRPLAFLNSV 289

DB 259 IYLVIIIVLTVFAVSIIPIHVMKTMNLRARLDFQTPAMCAFNDRVATYQVTRGLASLNSC 318

QY 290 INPVFYFLGDHFRDMLNQLR 311

DB 319 VDPILYFLAGDTFRRRLSRATR 340

RESULT 12

ID AAU10983 standard; Protein: 373 AA.

XX AAU10983;

XX 12-MAR-2002 (first entry)

XX Purinergic receptor P2Y, G-protein coupled 1.

XX Purinergic receptor P2Y, G-protein coupled 1; P2RY1; anticoagulant;

XX coagulant; platelet aggregation; haplotyping; drug screening;

XX transgenic animal; human.

XX Homo sapiens.

XX WO200190117-A2.

XX 29-NOV-2001.

XX 21-MAY-2001; 2001WO-US16432.

XX 19-MAY-2000; 2000US-205996P.

XX (GENA-) GENAISSANCE PHARM INC.

XX Kazemi A, Koshy B, Tanguay DA;

XX WPI, 2002-083074/11.

XX DR N-PSDB; AAS18599.

PT New purinergic receptor P2Y G-protein coupled 1 (P2RY1) gene  
PT polymorphic variants, useful e.g. in studying the expression and  
PT function of P2RY1 and screening candidate drugs for treating diseases  
PT related to P2RY1 activity -

PS Claim 28; Fig 3; 79pp; English.

CC The invention relates to a novel isolated polypeptide comprising a  
CC sequence which is a polymorphic variant of a reference sequence for the  
CC purinergic receptor P2Y, G-protein coupled, 1 (P2RY1) protein or its  
CC fragment. The polymorphic variant comprises one or more variant amino  
CC acids selected from valine at a position 34 and glycine at a position  
CC 262. The polymorphic variants are useful in studying the expression  
CC and function of P2RY1, in expressing P2RY1 protein for use in screening  
CC for candidate drugs to treat diseases related to P2RY1 activity, in  
CC studying the effect of the variation on the biological activity of  
CC P2RY1, and the binding affinity of candidate drugs targeting P2RY1 for  
CC the treatment of disorders related to platelet aggregation. The  
CC haplotyping methods are useful in validating P2RY1 as a candidate  
CC target for treating a specific condition or disease predicted to be  
CC associated with P2RY1 activity, or in the design of clinical trials of  
CC candidate drugs for treating a specific condition or disease associated  
CC with P2RY1 activity. The transgenic animals are useful for studying  
CC expression of the P2RY1 isogenes in vivo, for in vivo screening and  
CC testing of drugs targeted against P2RY1 protein, and for testing the  
CC efficacy of therapeutic agents and compounds for disorders related to  
CC platelet aggregation in a biological system. The present sequence  
CC represents the amino acid sequence of human purinergic receptor P2Y,  
CC G-coupled protein 1 (P2RY1).

XX Sequence 373 AA;

Query Match 28.5%; Score 497.5; DB 23; Length 373;

Best Local Similarity 36.3%; Pred. No. 5e-43; Mismatches 124; Indels 21; Gaps 9;

Matches 117; Conservative 60; Mismatches 124; Indels 21; Gaps 9;

QY 6 AW-NATCNMMLAA---BALEK---YLSIFYGIEFVGVGLNTTVVGYIFSLKNW 55

DB 24 SWGNSVASTAAVSSSFICALTKTGFOFYIPLPAVYILVFIIIGFLNSVAIMVFFHMKRW 83

QY 56 NSSNIYLFNLSVSDIAFLCTLPMLIRSYAN-GNWIYGDVLCISNRVYLANLYTSILFLT 114

DB 84 SGISVYMFNMLADFLVLTLPALIFYYFNKTDWIFGDMCKLQRFIFHVNLVGSILFLT 143

QY 115 FISIDRYLIKYPREHLLQKKEFAILISLAIVWVLTLELPILPLINPVITDNGT-TCN 173

DB 144 CISAHRYSGVYVPLKSLRLKKKNAICISVLWLVVAISILYSGTGVRKNNTITCY 203

QY 174 DPASGDPNVLIIYSMCLTLGLFIPFVM--CFPYVKIALFLKQRNRQVATAPL-EKP 230

DB 204 DTSDEYLRSYFIYSMCTTVAMFCVPLVILIGCYGLIVRALYKDLNLS---PLRRKS 258

QY 231 LNLVIMAVIFSVPPTPYHVMNRVRIASRLGSKQYOCT-QVINSFYIVTRPLAFLNSV 289

DB 259 IYLVIIIVLTVFAVSIIPIHVMKTMNLRARLDFQTPAMCAFNDRVATYQVTRGLASLNSC 318

QY 290 INPVFYFLGDHFRDMLNQLR 311

DB 319 VDPILYFLAGDTFRRRLSRATR 340

RESULT 13

ID AAU10985 standard; Protein: 373 AA.

XX AAU10985;

XX 12-MAR-2002 (first entry)

XX Purinergic receptor P2Y, G-protein coupled 1, isoform #2.

XX Purinergic receptor P2Y, G-protein coupled 1; P2RY1; anticoagulant;

XX coagulant; platelet aggregation; haplotyping; drug screening;



CC inverse agonists or partial agonists having applicability as therapeutic  
CC agents for treating diseases related to GPCR, e.g. lung cancer.  
CC Non-endogenous version of human GPCRs are also utilized in research  
CC settings and in vitro and in vivo system. Incorporating GPCRs can be  
CC utilized to elucidate and understand the roles these receptors  
CC play in the human condition, both normal and diseased.

XX  
XX Sequence 337 AA;  
SQ

Query Match 27.3%; Score 477; DB 22; Length 337;  
Best Local Similarity 35.9%; Pred. No. 5.9e-41;  
Matches 110; Conservative 61; Mismatches 123; Indels 12; Gaps 6;

QY 8 NATCKNMLAABALEKXYLSIFGYIEFVGVGNTIVVGYIFSLKNMNSNIYLFNLSV 67  
DB 23 NCTDEN-----IPLKMHYLPVITYGIIFLVGFPGNAVISTYIFKRPKMSSTIMMLAC 77  
QY 68 SDLAFLCTLPMLIRSYANG-NWYGDVLCISNRVYLANLYTSILFLPISIDRYLIKY 126  
DB 78 TDLVLTSLPFLIHYVASEGNNWIFGDFMCKFRFSPHFLYSILFLTCFSIFRYCVIHH 137  
QY 127 PPREHLQKKEPAIILSLAIWLVLTLELLPLPLINPVITDNGTGNDFASSGDPVNYLI 186  
DB 138 PMSCSIHKTRCAVAVACAVWMIISLVAVIPMTFLITSTRTNRSACLDITSSDELNTIKW 197  
QY 187 YSMCLTLGLFLPLFVWCFYFYKIALFLKQNRQVATLPLEKPLNLVIMAVIFSVPFT 246  
DB 198 YNLITATTFCPLVIVLVCYTTI-IHTLTGLOTDSCIK-QKARRLTILLLAFVYCL 255  
QY 247 PYHVRNRVRIASRLGSMKOYQCT-OVINSFYIVTRPLAFNSVINPVYFPLGDHFRDM 305  
DB 256 PFHILRVIRIESRLS---ISCSINQIHEAVYISRLAALNTFGLLLYVVSDNFQQA 312  
QY 306 LMNQLR 311  
DB 313 VCSIVR 318

RESULT 15  
AAOI5399  
ID AAOI5399 standard; protein, 337 AA.  
XX  
XX AAOI5399;  
AC  
XX  
XX 27-SEP-2002 (first entry)  
DT  
XX  
XX Human G protein-coupled receptor.  
DE  
XX  
XX Human; gene therapy; G protein-coupled receptor; drug development;  
KW central nervous system disease; endocrine disease; metabolic disease;  
KW cancer; respiratory disease; digestive disease; immune disease;  
KW inflammation; infection; circulatory disease.  
XX  
XX Homo sapiens.  
OS  
XX  
XX WO200257441-A1.  
PN  
XX  
XX 25-JUL-2002.  
PD  
XX  
XX 17-JAN-2002; 2002WO-JP00270.  
PF  
XX  
XX 18-JAN-2001; 2001JP-0010714.  
PR  
XX  
XX 30-MAR-2001; 2001JP-0102484.  
PR  
XX  
XX (TAKE ) TAKEDA CHEM IND LTD.  
PA  
XX  
XX Miwa M, Ito T, Shintani Y, Miyajima N;  
PI  
XX  
XX WPI; 2002-566800/60.  
DR  
XX  
XX N-PSDB; AAL43942.  
DR  
XX  
XX Human kidney-originated G protein-coupled receptor protein TGR30 and  
PT encoded DNA, for developing drugs to treat central nervous diseases,

PT endocrine diseases, metabolic diseases and cancer, including gene  
PT therapy -  
XX  
XX  
XX Claim 1; Page 88-90; 98pp; Japanese.  
PS  
XX  
XX The invention comprises the amino acid and coding sequence of a human G  
CC protein-coupled receptor. The DNA and protein sequences of the invention  
CC are useful for developing drugs to prevent or treat (gene therapy):  
CC central nervous system diseases; endocrine diseases; metabolic diseases;  
CC cancer; respiratory diseases; digestive diseases; immune diseases;  
CC inflammations; infections; and circulatory diseases. The present amino  
CC acid sequence represents the human G protein-coupled receptor of the  
CC invention.

XX  
XX Sequence 337 AA;  
SQ

Query Match 27.3%; Score 477; DB 23; Length 337;  
Best Local Similarity 35.9%; Pred. No. 5.9e-41;  
Matches 110; Conservative 61; Mismatches 123; Indels 12; Gaps 6;

QY 8 NATCKNMLAABALEKXYLSIFGYIEFVGVGNTIVVGYIFSLKNMNSNIYLFNLSV 67  
DB 23 NCTDEN-----IPLKMHYLPVITYGIIFLVGFPGNAVISTYIFKRPKMSSTIMMLAC 77  
QY 68 SDLAFLCTLPMLIRSYANG-NWYGDVLCISNRVYLANLYTSILFLPISIDRYLIKY 126  
DB 78 TDLVLTSLPFLIHYVASEGNNWIFGDFMCKFRFSPHFLYSILFLTCFSIFRYCVIHH 137  
QY 127 PPREHLQKKEPAIILSLAIWLVLTLELLPLPLINPVITDNGTGNDFASSGDPVNYLI 186  
DB 138 PMSCSIHKTRCAVAVACAVWMIISLVAVIPMTFLITSTRTNRSACLDITSSDELNTIKW 197  
QY 187 YSMCLTLGLFLPLFVWCFYFYKIALFLKQNRQVATLPLEKPLNLVIMAVIFSVPFT 246  
DB 198 YNLITATTFCPLVIVLVCYTTI-IHTLTGLOTDSCIK-QKARRLTILLLAFVYCL 255  
QY 247 PYHVRNRVRIASRLGSMKOYQCT-OVINSFYIVTRPLAFNSVINPVYFPLGDHFRDM 305  
DB 256 PFHILRVIRIESRLS---ISCSINQIHEAVYISRLAALNTFGLLLYVVSDNFQQA 312  
QY 306 LMNQLR 311  
DB 313 VCSIVR 318

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Job time : 40 secs

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